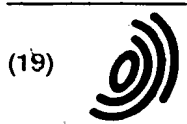


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Europäisches Patentamt

(19)

European Patent Office

Office européen des brevets



(11)

EP 0 763 522 A1

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication:

19.03.1997 Bulletin 1997/12

(51) Int. Cl.⁶: **C07C 233/47**, C08F 26/02

(21) Application number: **96114552.1**

(22) Date of filing: **11.09.1996**

(84) Designated Contracting States:

DE FR GB

(30) Priority: **13.09.1995 US 527311**

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(54) **N-vinylformamide/alkyl acrylate michael adducts**

(57) Novel unsaturated compounds comprising the 3-(N-vinylformamido)propionates and 2-methyl-3-(N-vinylformamido)propionates, or the corresponding propionamides, are obtained by the Michael addition reaction of N-vinylformamide with an acrylic or methacrylic acid ester. These compounds are useful as monomers in free radical polymerizations, particularly as components of photocurable coatings.

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Description

CROSS-REFERENCE TO PARENT APPLICATION

5 This is a continuation-in-part of copending application serial no. 08/246,434, the subject matter of which is hereby incorporated by reference.

TECHNICAL FIELD OF THE INVENTION

10 The invention relates to compounds useful as monomers in free radical polymerizations, particularly as components of photocurable coatings.

BACKGROUND OF THE INVENTION

15 Unsaturated monomers of the N-vinylamide class have been employed in free radical polymerization reactions for the preparation of homopolymers and copolymers having a range of useful properties. Of these, the N-vinyl lactams, in particular N-vinyl-2-pyrrolidone, have been employed extensively owing to their wide commercial availability and rapid polymerization with a range of comonomers including the acrylic and methacrylic esters. Polymerizations of other cyclic and acyclic N-alkyl substituted (tertiary) N-vinyl carboxylic acid amides have been described as well. Secondary N-vinyl-
20 lamides such as N-vinylformamide and N-vinylacetamide also copolymerize well with acrylic compounds and are useful for preparing hydrophilic polymers which may be subsequently hydrolyzed to introduce vinylamine functionality.

Several N-vinylamide monomers have been suggested as components of liquid and solid photopolymerizable compositions of various types. As a class, the vinylamides possess certain useful properties for these applications, including good compatibility with other chemistries, copolymerizability with acrylates, comparative resistance to oxygen
25 inhibition, and favorable adhesion-promoting characteristics.

Lorenz et al. (U.S. Patent 4,129,709) disclose a coating composition comprising N-vinyl-2-pyrrolidone, an acrylated oligomer, and an acrylic acid ester having a boiling point of at least 200°C at 760mm Hg. These compositions may be cured by exposure to actinic radiation between 200 and 750 nm or by an electron beam. Tu et al. (U.S. Pat. 4,319,811) describe radiation curable coatings consisting of triacrylate or tetraacrylate monomers with an N-vinyl imido monomer, preferably an N-vinyl lactam such as N-vinyl-2-pyrrolidone. Priola and coworkers (U.S. Pat. 4,348,427) describe compositions comprising mixtures of acrylated oligomers and/or unsaturated polyester oligomers with a least one unsaturated compound of the amide, lactam, piperidone and urea classes, and curing them by exposure to ultraviolet radiation in the 200-400 nm range.
30

Cornforth et al. (U.S. Pat. 5,281,682) teach improved radiation-curable formulations containing N-vinylformamide and an oligomer selected from the group epoxy acrylates, urethane acrylates, polyester acrylates and mixtures thereof. Elzer et al. (U.S. Pat. 4,725,524) disclose a dry film photoresist containing an acrylic or methacrylic oligomer, a compatible film-forming water-soluble polymer, one or more compatible photopolymerizable monomers, a photoinitiator, and other additives. Barzynski et al. (U.S. Pat. 4,205,139 and 4,424,314) teach curable compositions containing N-vinyl compounds in which at least two N-vinyl groups are present and in which at least one carbonyl group is bound to the nitrogen of the N-vinyl group, said carbonyl group in turn being bonded to a nitrogen or carbon atom.
40

The most commercially important class of radiation-curable compositions relies upon the free radical photopolymerization of unsaturated acrylic compounds. These systems are commonly based on acrylic ester-terminated oligomers derived from one of several major resin chemistries, such as urethanes, epoxies, polyesters, and others. The acrylated oligomers are often compounded with various nonpolymerizable materials (pigments, fillers, flow agents, antioxidants, etc), as well as photoinitiators and co-catalysts, and applied to a substrate before curing. Curing is accomplished by exposing the formulation to ultraviolet light or other type of radiation until a dry adherent polymerized film is formed. Formulations of this general description find use as printing inks, protective coatings, adhesives and the like.
45

In practice, it is often necessary to incorporate diluent monomers into these formulations in order to lower the viscosity of the oligomers sufficiently to permit adequate flow and leveling on the substrate prior to irradiation. This is particularly true of formulations applied by methods common to the printing and coating industries. Diluent monomers used for this purpose fall into two broad categories: multifunctional (or polyunsaturated) types and monofunctional (or monounsaturated) compounds. Multifunctional monomers generally provide high cure speed and high crosslink density, leading to hard, chemically-resistant films, but they may not sufficiently lower viscosity and may also contribute to poor adhesion due to excessive shrinkage of the films on curing. Monofunctional monomers are usually more effective in reducing viscosity, and can yield softer, more extensible films that exhibit better adhesion to many substrates.
50

55 Although a large number of unsaturated compounds are potentially useful as diluents, relatively few have proven to be practical. This is because reactive monomers employed in this technology must conform to a number of physical and performance requirements, among them: high fluidity, low vapor pressure, low color, low odor, high flash point, low toxicity and irritancy, broad compatibility and rapid copolymerizability with the other unsaturated species.

Monounsaturated N-vinylamides have the aforementioned useful properties as photocurable compounds, however, few of these monomers meet all the criteria for use as diluents. It is found, for example, that the lower alkyl tertiary N-vinylamides (e.g. N-methyl-N-vinylformamide, N-methyl-N-vinylacetamide) are excellent solvents, but are excessively volatile and possess relatively low flash points (under 200°F). The higher secondary and tertiary N-vinylamides (e.g. N-vinylacetamide, N-vinylbenzamide, N-vinylcaprolactam, N-phenyl-N-vinylacetamide, N-vinyl-2-piperidinone, N-vinyl-succinimide, N-vinylmorpholinone) are less volatile and less hydrophilic, but have melting points above room temperature, making them less desirable as diluents or coreactants in liquid systems. Many of these monomers exhibit unacceptably slow cure rates as well.

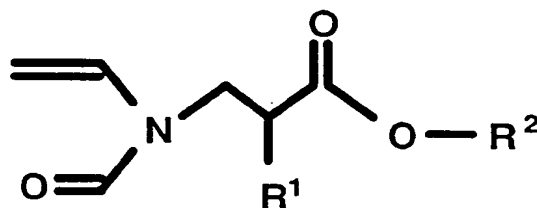
Kurtz and Disselnkotter (*Liebigs Ann. Chem.*, 764, p. 69-93, 1972) describe the preparation of 3-N-vinylformamido propionitrile by the reaction of N-vinylformamide with acrylonitrile in benzene utilizing a potassium cyanide catalyst. Applications for the compound are not reported. However, this material has also been found to have certain deficiencies as a photocurable monomer. A further disadvantage is the use of a highly toxic catalyst in its preparation.

Of the N-vinyl class, only N-vinyl-2-pyrrolidone (NVP) has achieved widespread commercial acceptance in the technology owing to its satisfactory combination of good solvent properties, low vapor pressure and high cure rate in liquid photocure systems. More recently, N-vinylformamide (NVF) has been shown to offer equal or superior performance. However, both NVP and NVF have certain disadvantages as well. Notably, both monomers possess relatively high glass transition temperatures (150 - 175°C) which imposes some constraints upon the film properties that can be achieved. In addition, both monomers are relatively hydrophilic which can undesirably increase the water sensitivity of cured films.

It is the object of this invention to provide novel N-vinyl monomers which simultaneously exhibit the desirable solvent and polymerization characteristics of the N-vinylamides, and which also possess the required physical properties necessary to render them practical as components of liquid radiation-curable formulations, and which further possess a wider range of Tg and hydrophobic properties than is exhibited by the preferred liquid monomers of the prior art, i.e. NVP and NVF.

SUMMARY OF THE INVENTION

The present invention relates to unsaturated monomers consisting of the 3-(N-vinylformamido)propionates and 2-methyl-3-(N-vinylformamido)propionates, or the corresponding propionamides, obtained from the Michael addition of N-vinylformamide (NVF) to acrylic or methacrylic acid esters. The materials encompassed by this invention have the general structure:



Where R¹ is hydrogen or methyl, and R² is a linear or branched alkyl containing from 9 to 20 carbon atoms, a cyclic or polycyclic alkyl, halogenated alkyl, arylalkyl, or alkylene group, each containing up to about 20 carbon atoms.

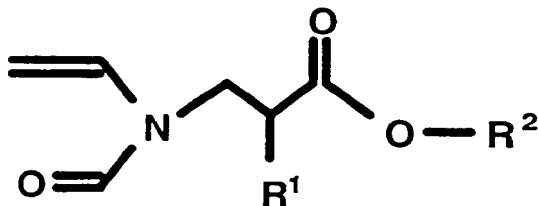
The compounds of the instant invention are prepared by the reaction of N-vinylformamide with the acrylic or methacrylic ester in the presence of a basic catalyst, such as sodium methoxide, sodium ethoxide, or butyllithium. The reaction is performed either in a mixture of the neat monomers, or in a suitable organic solvent, at a temperature between 0 and 100°C. The reaction is allowed to proceed for a period between 0.1 and 100 hours, after which time the purified product is preferably recovered by vacuum distillation.

Compounds prepared according to this invention have been found to be useful as monomers, especially in photocurable coatings. The compounds demonstrate excellent solvent properties, low viscosity, a wide liquid range, low vapor pressure, low color, excellent color stability, good adhesion and high cure speed in free radical photopolymerization. In addition, the materials defined under the invention may be selected to provide a wide range of Tg and hydrophobic characteristics.

It is recognized by the inventors that, while useful in radiation curing, these monomers may have broader utility in other types of polymer synthesis such as, for example, bulk, suspension, emulsion or solution polymerization, and with comonomers different from those commonly used in radiation-curing (e.g. ethylene, vinyl acetate, vinyl chloride, maleate esters, acrylonitrile, etc.). Therefore, our invention is intended to subsume the homopolymerization or copolymerization of these monomers by any means known in the art.

DETAILED DESCRIPTION OF THE INVENTION

Novel unsaturated monomers consisting of the 3-(N-vinylformamido)propionates and 2-methyl-3-(N-vinylformamido)propionates obtained from the Michael addition of N-vinylformamide to an acrylic or methacrylic acid ester. The materials encompassed by this invention have the general structure:



where R^1 is hydrogen or methyl, and R^2 is a linear or branched alkyl containing from 9 to 20 carbon atoms, a cyclic or polycyclic alkyl, halogenated alkyl, arylalkyl, or alkylene group, each containing up to about 20 carbon atoms.

The subject compounds are readily prepared by the nucleophilic addition of N-vinylformamide (NVF) to an acrylic or methacrylic acid ester. The reaction can be carried out in a simple mixture of the neat monomers, or in a solvent. NVF and the (meth)acrylate ester groups are present in the reaction mixture in molar ratio of from about 1:10 to about 20:1, and preferably at a ratio of about 1.1 moles NVF per equivalent of (meth)acrylate. A free radical inhibitor, such as benzoquinone, is also added. The reaction is preferably carried out in air, at atmospheric or other pressure.

The reaction is conducted in the presence of a strongly basic catalyst such as the alkali metal or quaternary amine hydroxides or alkoxides. Bases of the methoxide, ethoxide, isopropoxide and t-butoxide class are preferred. Sodium methoxide is especially preferred. Other useful catalysts include the aryl- and alkylolithiums, potassiums and sodiums. The use of other strong bases, such as tertiary amines, amidines or metal amides, such as sodium amide or lithium diisobutylamide is also conceivable, as is the use of anhydrous carbonates. The catalyst is present in the reaction mixture in an amount between about 0.0005 and about 5 wt.%, and preferably between 0.1 and 0.5 wt.% of the total weight of the reactants. In general, the appropriate level of catalyst will vary depending upon the equivalent weight of the (meth)acrylic ester and the molar ratio of the reactants. Following the addition of catalyst, the reactants or reactant solution are maintained at a temperature between 0 and 100°C, and preferably between 20 and 60°C, and allowed to react for a period of from about 0.1 hours to about 100 hours.

In principle, the reagent monomers can be reacted in batch fashion, via staged addition, or continuously, whichever is most suitable. Synthesis is advantageously performed in a mixture of the neat monomers, however, an inert solvent for both reactants may be employed. Potential solvents include the amides, lower hydrocarbons, chlorinated hydrocarbons, and aromatics. Preferred solvents are esters (e.g. ethyl acetate) and ethers (e.g. ethyl ether, tetrahydrofuran).

The purified 3-(N-vinylformamido)propionate adduct is preferably recovered from the reaction mixture by distillation under vacuum at between 0.01 and about 50 torr. Distillation may be performed either batchwise or continuously as, for example, in a wiped film evaporator. Other possible methods of purification include neutralization of the catalyst followed by vacuum stripping of the unreacted starting materials, or solvent extraction. In some cases, isolation of the 3-N-vinylformamido adduct from the reaction mixture may not be required. For example, the crude product may be suitable for applications in which quantities of unreacted NVF and/or (meth)acrylic ester are not problematic.

A wide range of acrylic and methacrylic monomers are potentially useful as the coreactant or Michael acceptor in this technology. The most suitable Michael acceptors are the (meth)acrylic monoesters which do not contain a source of active hydrogen in their structure. Such undesirable sources of active hydrogen are, for example, O-H and N-H species present in alcohol and carboxylic acid groups, primary and secondary amines, primary and secondary amides, etc.

Notwithstanding this provision, the nature of the (meth)acrylic ester group can be selected from a wide variety of structures ranging in size from one to about twenty carbon atoms, including linear and branched alkyl, cyclic and polycyclic alkyl, arylalkyl, tertiary aminoalkyl, unsaturated alkyl, halogenated alkyl, etc. Examples of suitable (meth)acrylic esters for reaction with NVF include methyl acrylate, ethyl acrylate, butyl acrylate, ethylhexyl acrylate, isooctyl acrylate, isobornyl acrylate, benzyl acrylate, N,N-dimethylaminoethyl acrylate, allyl acrylate and methyl methacrylate.

It will be apparent to those skilled in the art that the (meth)acrylate esters above span a range of properties, such as T_g and hydrophobicity. It has been found that by adducting NVF to an appropriate (meth)acrylic ester, the T_g and hydrophobic properties of the Michael acceptor (i.e. acrylate or methacrylate ester) can be incorporated into the resulting N-vinylamide monomer. Further, it has been found that acrylate esters which are normally too volatile or too flammable for use in radiation curing (e.g. methyl acrylate, butyl acrylate, and ethylhexyl acrylate) can be employed as the Michael acceptor for NVF to yield 3-N-vinylformamido adducts that exhibit low volatility and high flash points. The table below compares the physical properties of some 3-(N-vinylformamido)propionates prepared under this invention with

N-vinylamides of the prior art.

Compound	Appearance (25°C)	Boiling point	Flash point (Cel.)	Tg (homopoly.)
Methyl 3-(N-vinylformamido)-propionate	Colorless liquid	75°C/0.8 mmHg	124 ¹	24 °C
Ethyl 3-(N-vinylformamido)-propionate	Colorless liquid	88°C /1 mmHg	124 ¹	46 °C
Butyl 3-(N-vinylformamido)-propionate	Colorless liquid	96°C / 0.5 mmHg	98 ¹	---
t-Butyl 3-(N-vinylformamido)-propionate	Colorless liquid	80°C/0.5 mmHg	-----	-----
Ethylhexyl 3-(N-vinylformamido)-propionate	Colorless liquid	122°C/0.8 mmHg	---	---
Methyl 2-methyl-3-(N-vinylformamido)-propionate	Colorless liquid	76°C/0.5 mmHg	---	---
N-methyl-N-vinylacetamide	Yellow liquid	70°C/25 mmHg	58	---
N-vinylformamide	Sl. yellow liquid.	84°C/10 mmHg	102	150 °C
N-vinyl-2-pyrrolidone	Sl. yellow liquid.	96°C/14 mmHg	98	175 °C
3-(N-vinylformamido)-propionitrile	Yellow liquid	99°C/1.5 mmHg	~150	---

¹ASTM D-3828

As indicated above, the 3-(N-vinylformamido)propionate esters are colorless liquids at room temperature, exhibit advantageously low volatility, and possess flash points equivalent to or higher than NVF and NVP. Moreover, as evidenced by the examples of the methyl and ethyl propionate esters, adduction of NVF to the corresponding acrylate yields an N-vinyl monomer with a significantly lower Tg than the N-vinylformamide starting material.

For comparison, the NVF/acrylonitrile adduct (3-(N-vinylformamido)propionitrile) of the prior art was prepared according to the method of Kurtz and Disselinkotter. The distilled product was found to be colored and to exhibit poor color stability. Moreover, the color of the propionitrile monomer was found to persist in cured films containing the material. Such coloration is regarded as a disadvantage in this technology since it could interfere with the spectral characteristics of pigments or dyes which may be incorporated into the formulations, as for example, when they are employed as tinted coatings or inks, or where the formulation is intended to act as a clear varnish over a colored background or graphic. By contrast, the propionate ester adducts of the present invention are essentially colorless as produced and demonstrate excellent resistance to discoloration even after protracted storage at elevated temperatures. A further advantage of the propionate esters is their lower neat viscosity which, as will be shown in the following examples, results in their greater effectiveness in reducing the viscosity of acrylated oligomers.

Compound	Viscosity (25°C)	Gardner Color (initial)	Gardner Color (20 days at - 20°C)
Butyl 3-(N-vinylformamido)-propionate	13.5 cps	<1	<1
3-(N-vinylformamido)-propionitrile	39.1 cps	5 - 6	6 - 7

In addition to their use in radiation-curable formulations, The 3-(N-vinylformamido)-propionate esters will be useful in conventional polymer synthesis for the preparation of water- or organic solvent-soluble homopolymers and copolymers which may have utility in various commercial applications. In the case of copolymers, the propionate esters of the present invention can be copolymerized with one or more copolymerizable monomers selected from the group consisting of vinyl esters, vinylamides, acrylic esters, methacrylic esters, maleic esters, substituted and unsubstituted acrylamides, substituted and unsubstituted methacrylamides, diallyl monomers, maleic acid, maleic anhydride, ethylene, vinyl chloride, acrylonitrile, cationic acid and acrylic acid. Potential uses for such derived polymers include the following: coating resins for metal, plastics and paper; adhesives; hair conditioning and fixative resins; paper strength and sizing agents; water treatment chemicals; pigment dispersants and binders for printing inks; colloids for emulsion polymeriza-

tion; hydrogels.

The following examples are intended to better illustrate the invention and are not meant to be limiting.

EXAMPLE 1

Preparation of methyl 3-(N-vinylformamido)propionate

To a 1000 mL three-neck round bottom flask equipped with a cold water condenser and stirrer was added 215 grams (2.5 moles) of methyl acrylate, 195 grams (2.75 moles) of N-vinylformamide, and 0.1 gram of benzoquinone. The mixture was stirred at ambient temperature for two minutes and 1.5 grams sodium methoxide was added in one portion. The mixture was stirred for approximately 2 hours at ambient temperature and allowed to stand overnight. The reaction mixture was distilled under vacuum and 361 grams of product (92% yield) boiling at 75°C (0.8mm Hg) was collected for analysis. ¹H NMR analysis confirmed the identity of the 3-N-vinylformamido adduct.

EXAMPLE 2

Preparation of ethyl 3-(N-vinylformamido)propionate

To a 100 mL three-neck round bottom flask equipped with a cold water condenser and stirrer was added 30.9 grams of ethyl acrylate and 21.9 grams of NVF. After mixing for 2 minutes, 0.20 grams of sodium methoxide was added in one portion. The mixture was allowed to react for 3 hours and then distilled to recover 40.3 grams of product boiling at 88°C (1.0 mm Hg).

EXAMPLE 3

Preparation of butyl 3-(N-vinylformamido)propionate

72.5 grams of N-butyl acrylate were reacted with 43.0 grams of NVF according to the procedure of Example 2. The mixture was allowed to react for 4 hours and then distilled to recover 101.2 grams of product boiling at 96°C (0.5 mm Hg).

EXAMPLE 4

Preparation of t-butyl 3-(N-vinylformamido)propionate

21.9 grams of t-butyl acrylate were reacted with 42.0 grams of NVF according to the procedure of Example 2. The mixture was allowed to react for 72 hours and then distilled to recover 51.5 grams of product boiling at 80°C (0.5 mm Hg).

EXAMPLE 5

Preparation of ethylhexyl 3-(N-vinylformamido)propionate

83.1 grams of 2-ethylhexyl acrylate were reacted with 34.4 grams of NVF according to the procedure of Example 2. The mixture was allowed to react for 96 hours and then distilled to recover 79.8 grams of product boiling at 122°C (0.8 mm Hg).

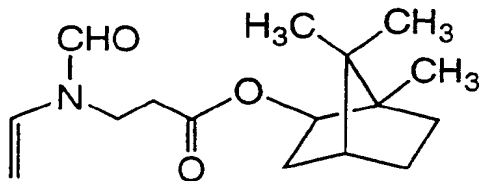
EXAMPLE 6

Preparation of methyl 2-methyl-3-(N-vinylformamido)propionate

To a 100 mL three neck round bottom flask equipped with a cold water condenser, oil bath and stirrer was added 33.2 grams of methyl methacrylate and 23.6 grams of NVF. The mixture was stirred at room temperature for 2 minutes and 80 mg of butyllithium (as a 2.5M solution in hexane) was added in a single portion. The reaction mixture was agitated for 8 hours at 65°C and then distilled to recover 29.5 grams of product boiling at 75-77°C (0.5 mm Hg).

EXAMPLE 7**Cycloalkyl acrylate:**

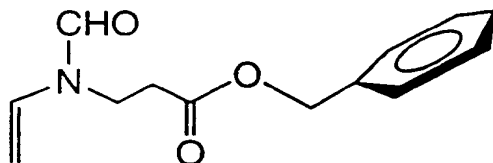
5 Preparation of isobornyl 3-(N-vinylformamido)propionate



15 To a 100 mL three-neck round bottom flask equipped with a cold water condenser and stirrer was added 20.8 grams of isobornyl acrylate, 0.01 gram of benzoquinone, and 7.1 grams of NVF. After mixing for 2 minutes, 0.1 gram of 25% sodium methoxide methanol solution was added in one portion. After stirring at ambient temperature for 20 hours, the colored mixture was distilled and a fraction boiling at 143-145°C/0.7 mmHg was collected to give 22.3 grams of colorless isobornyl 3-(N-vinylformamido)propionate (80%).

EXAMPLE 8**Aromatic moiety containing acrylate:**

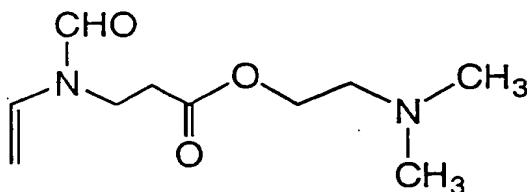
25 Preparation of benzyl 3-(N-vinylformamido)propionate



35 To a 100 mL three-neck round bottom flask equipped with a cold water condenser and stirrer was added 14.6 grams of benzyl acrylate, 0.01 gram of benzoquinone, and 7.1 grams of NVF. After mixing for 2 minutes, 0.1 gram of 25% sodium methoxide methanol solution was added in one portion. After stirring at ambient temperature for 24 hours, the colored mixture was distilled and a fraction boiling at 147-148°C/0.8 mmHg was collected to yield 16.5 grams of benzyl 3-(N-vinylformamido)propionate as a pale yellow liquid (76%).

EXAMPLE 9**Nitrogen containing acrylate:**

45 Preparation of N,N-dimethylaminoethyl 3-(N-vinylformamido)propionate



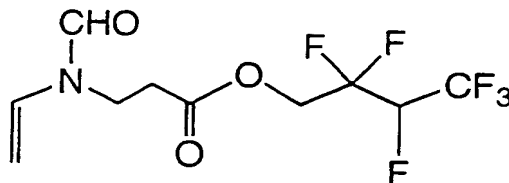
55 To a 100 mL three-neck round bottom flask equipped with a cold water condenser and stirrer was added 14.3 grams of N,N-dimethylaminoethyl acrylate, 0.01 gram of benzoquinone, and 7.1 grams of NVF. After mixing for 2 minutes, 0.1 gram of 25% sodium methoxide methanol solution was added in one portion. After stirring at ambient temper-

ature for 24 hours, the color mixture was distilled and a fraction boiling at 115-120°C/1 mmHg was collected to yield 12.2 grams of N,N-dimethylaminoethyl 3-(N-vinylformamido)propionate as a pale yellow liquid (57%).

EXAMPLE 10

Fluoroalkyl acrylate:

Preparation of 2,2,3,4,4,4-Hexafluorobutyl 3-(N-vinylformamido)propionate

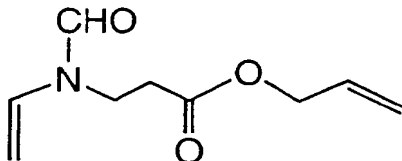


To a 100 mL three-neck round bottom flask equipped with a cold water condenser and stirrer was added 23.6 grams of 2,2,3,4,4,4-hexafluorobutyl acrylate, 0.01 gram of benzoquinone, and 7.1 grams of NVF. After mixing for 2 minutes, 0.2 gram of 25% sodium methoxide methanol solution was added in one portion. After stirring at ambient temperature for 24 hours, the colored mixture was distilled and a fraction boiling at 95°C/0.5 mmHg was collected to yield 14.3 grams of 2,2,3,4,4,4-hexafluorobutyl 3-(N-vinylformamido)propionate as a pale yellow liquid (47 %).

EXAMPLE 11

Allyl acrylate:

Preparation of Allyl 3-(N-vinylformamido)propionate



To a 250 mL three-neck round bottom flask equipped with a cold water condenser and stirrer was added 82.2 grams of allyl acrylate, 0.01 gram of benzoquinone, and 57.3 grams of NVF. After mixing for 2 minutes, 0.2 gram of 25% sodium methoxide methanol solution was added in one portion. After stirring at 60°C for 24 hours, the colored mixture was distilled and a fraction boiling at 100-110°C/1 mmHg was collected to yield 53.0 grams of allyl 3-(N-vinylformamido)propionate as a pale yellow liquid (38 %).

EXAMPLE 12

Homopolymerization of methyl 3-(N-vinylformamido)propionate

The homopolymer of the NVF/methyl acrylate adduct was prepared in order to measure the glass transition temperature of the material. 50 grams of the adduct prepared in Example 1 were added to a reaction vessel, equipped with a stirrer and oil bath, and containing 60 grams of toluene. The mixture was agitated and the temperature raised to 100°C and 0.5 grams of Vazo™ 88 initiator (1,1'-azobiscyclohexanecarbonitrile) was added in one portion. The reaction was allowed to proceed for 32 hours after which the precipitated polymer was recovered by filtration. The homopolymer was analyzed by gel permeation chromatography and found to have a number average molecular weight (Mn) of 55,600 and a weight average molecular weight (Mw) of 152,100. The glass transition temperature was measured using a differential scanning calorimeter (DSC) and found to be 24°C.

EXAMPLE 13**Homopolymerization of ethyl 3-(N-vinylformamido)propionate**

5 The homopolymer of NVF/ethyl acrylate adduct was also prepared. 15.1 grams of the adduct prepared in Example 2 was added to a reaction vessel, equipped with a stirrer and oil bath, into which had been placed 13 grams of ethyl acetate. The mixture was agitated and the temperature raised to 80°C, and 0.03 grams of Vazo™ 67 (2,2'-azobis-2-methylbutanenitrile) initiator was added in one portion. The reaction was allowed to proceed for 29 hours after which the polymer was recovered by precipitation into water, filtered and dried in a vacuum oven. Analysis by gel permeation chromatography indicated a Mn of 32,300 and a Mw of 106,900. The glass transition temperature as measured by DSC was found to be 46°C.

EXAMPLE 14**15 Copolymerization of methyl 3-(N-vinylformamido)propionate (NVF/MA) with NVF**

A three neck round bottom flask equipped with a mechanical stirrer, a water condenser, nitrogen inlet/outlet tube and an addition funnel was charged with 15.4 g of NVF/MA, 0.2 g of Vazo™ 88, and 36.1 g of methoxypropanol. The mixture was heated to 100°C, and stirred at that temperature for 30 minutes. A mixture of 4.6 g of NVF and 0.2 g of Vazo™ 88 in 10.5 g of methoxypropanol was added through the addition funnel over 40 minutes. The mixture was heated continuously at 100°C for four hours. A sample withdrawn from the reaction solution was analyzed by GC, which indicated that 76% of the NVF/MA and 98% of the NVF were converted to polymer. The solvent was removed under vacuum and the product was dissolved in D₂O. ¹³C NMR analysis showed the expected copolymer with a 40/60 ratio of NVF to NVF/MA.

EXAMPLE 15**Copolymerization of methyl 3-(N-vinylformamido)propionate with butyl acrylate**

30 A three neck round bottom flask, equipped with a mechanical stirrer, a water condenser, nitrogen inlet/outlet tube and an addition funnel, was charged with 11.5 g of NVF/MA, 0.1 g of Triginox 23, and 20.4 g of methanol. The mixture was heated to 60°C and stirred at that temperature for 30 minutes. A mixture of 9.4 g of butyl acrylate and 0.1 g of Triginox 23 in 20.1 g of methanol was added through the addition funnel over 3 hours. The mixture was heated continuously at 60°C for two hours. A sample withdrawn from the reaction solution was analyzed by GC, which indicated that 77% of the NVF/MA and 100% of the butyl acrylate were converted to polymer. The solvent was removed under vacuum and the product was dissolved in DMSO. ¹³C NMR analysis showed the expected copolymer with a 53:47 ratio of butyl acrylate to NVF/MA. The molecular weight of this polymer, determined by GPC (polystyrene as standard), was: Mw = 105,485 and Mn = 33,616.

EXAMPLE 16**Copolymerization of methyl 3-(N-vinylformamido)propionate with vinyl propionate**

45 A three neck round bottom flask, equipped with a mechanical stirrer, a water condenser, nitrogen inlet/outlet tube and an addition funnel, was charged with 12.4 g of NVF/MA, 7.9 g of vinyl propionate, and 35.8 g of acetonitrile. The mixture was heated to 60°C, and the mixture was stirred at that temperature for 30 minutes. Triginox 23, 0.2 g, in 10.5 g of acetonitrile was added in one portion to the reaction flask. The mixture was heated continuously at 60°C for 47 hours. A sample withdrawn from the reaction solution was analyzed by GC, which indicated that 83% of the NVF/MA and 76% of the vinyl propionate were converted to polymer. The solvent was removed under vacuum and the product was dissolved in deuterioacetonitrile. ¹³C NMR analysis showed the expected copolymer with a 50:50 ratio of vinyl propionate to NVF/MA. The molecular weight of this polymer, determined by GPC (polystyrene as standard), was: Mw = 19,338 and Mn = 8313. The glass transition temperature of this polymer was about 40°C.

EXAMPLE 17

55 The relative performance of the NVF/acrylate ester adducts as diluents in photopolymerizable compositions was assessed in comparison with N-vinyl monomers of the prior art. Equal weight fractions of the N-vinyl compounds were compared in a model formulation containing an epoxy acrylate oligomer, two multifunctional acrylate monomers and a common free radical photoinitiator.

Component	Weight %
Epoxy diacrylate oligomer ¹	50%
TMPTA ²	10%
TRPGDA ³	10%
N-Vinyl Monomer	30%
Irgacure™ 184 ⁴	2.5 phr (based on weight of the above)

¹Ebecryl™ 3700 (UCB Radcure)

²Trimethylolpropane triacrylate (UCB Radcure)

³Tripropyleneglycol diacrylate (Sartomer SR-306)

⁴1-Hydroxycyclohexyl phenyl ketone (Ciba-Geigy)

The liquid mixtures comprising the oligomer, acrylate monomers, N-vinyl monomer, and photoinitiator were mixed well and the Brookfield viscosity was measured. Thin films were drawn down on cleaned 3"x 5" aluminum panels using a #10 wire bar. The panels were cured under ultraviolet light in air using a commercial 300 watt/inch medium pressure mercury lamp and conveyor system. Cured film properties were assessed after a single exposure at a conveyor speed of 105 feet per minute.

The extent of cure was indicated by measuring the water and solvent (methyl ethyl ketone) resistance of the films using the double rub test. Film hardness was evaluated by the Persoz hardness technique using a BYK Gardner Pendulum Hardness Tester calibrated on glass (412 seconds). Relative cure speed of each formulation was evaluated using a depth of cure comparator (UV Process Supply Company).

The properties of formulations containing three N-vinyl monomers of the prior art (N-vinylformamide, N-vinyl-2-pyrrolidone and N-methyl-N-vinylacetamide) are listed below:

	NVF	NVP	NMNVA
Flashpoint (°C)	102	98	58
Formulation viscosity (cps, 25°C)	625	464	212
Persoz hardness (seconds)	255	252	178
Depth of cure (mils)	Ca.90	Ca.90	Ca.30
Water double rubs	>200	>200	>200
MEK double rubs	>200	>200	>200
Color of cured film	Colorless	Colorless	Yellow

As expected, NVF and NVP produced hard, colorless, chemically resistant films, and also exhibited excellent cure speed in this system. NMNVA was most effective in reducing viscosity and produced softer films that retained the yellow color of the monomer. The cure depth of NMNVA was significantly less than the other monomers which is indicative of a slower rate of photopolymerization. In general, a high figure is desirable in this test since it implies that a shorter exposure time, or higher line speed, can be used to cure a given film thickness.

Formulations containing the 3-(N-vinylformamido)propionate esters were then evaluated according to the procedure above. Methyl 3-(N-vinylformamido)propionate (abbreviated below as NVF/MA), ethyl 3-(N-vinylformamido)propionate (NVF/EA), and n-butyl 3-(N-vinylformamido)propionate (NVF/BA) were formulated and cured alongside the 3-(N-vinylformamido)propionitrile (NVF/AN) compound of the prior art:

	NVF/MA	NVF/EA	NVF/BA	NVF/AN
Formulation viscosity (cps, 25°C)	314	1080	1250	1660
Persoz hardness (seconds)	159	233	160	219
Depth of cure (mils)	78	70	55	55
Water double rubs	>200	>200	>200	>200
MEK double rubs	>200	>200	>200	>200
Color of cured film	Colorless	Colorless	Colorless	Yellow

As shown, all three of the propionate esters were more effective as diluents than the propionitrile adduct. All of the NVF adducts cured to a dry surface in a single exposure to give chemically resistant crosslinked films. The NVF/methyl acrylate adduct gave a lower formulation viscosity than NVF and NVP, produced the softest films and showed excellent cure speed. All of the adducts were observed to reduce the hardness of the films relative to NVF, as would be expected based on their lower Tg values.

The NVF/acrylonitrile adduct produced cured films that retained a strong yellow color. These were measured according to the CIE scale and determined to have a delta-E value of 26 (strongly colored) versus a colorless control (delta-E of 0). All of the NVF/acrylate ester products cured to colorless films.

EXAMPLE 18

The water sensitivity of cured films containing methyl 3-(N-vinylformamido)-propionate (NVF/MA) was compared with an identical formulation containing NVF. The formulation used in this test was as follows:

Component	Weight %
Urethane diacrylate oligomer ¹	50%
TMPTA ²	10%
TRPGDA ³	10%
N-Vinyl Monomer	30%
Irgacure™ 184 ⁴	2.5 phr (based on weight of the above)

¹ Airthane™ 504 prepolymer capped with 2-hydroxyethyl acrylate (Air Products)

² Trimethylolpropane triacrylate (UCB Radcure)

³ Tripropyleneglycol diacrylate (Sartomer SR-306)

⁴ 1-Hydroxycyclohexyl phenyl ketone (Ciba-Geigy)

The formulations were cured under the conditions shown in Example 14 and the degree of water sorption of a weighed film was measured after immersion in water for one hour at 60°C. The soaked films were tamped dry with a paper towel, reweighed, and then dried in an oven at 100°C for two hours. The final weight of the film after oven drying was also recorded. The swell index is defined as the weight ratio of the soaked film to the original starting weight. The % soluble in water is defined as the decrease in weight after oven drying compared to the starting weight. The results of this test are shown below:

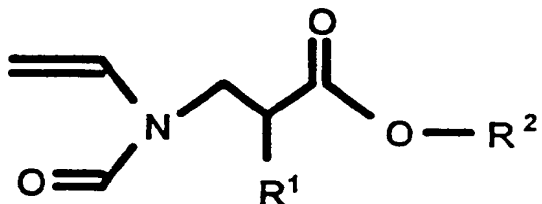
	NVF	NVF/MA
Persoz hardness (seconds)	308	167
Water double rubs	>200	>200
MEK double rubs	>200	>200
Swell index (water)	1.2	1.0
% Soluble	7%	7%

As shown, the cured film containing methyl 3-(N-vinylformamido)propionate exhibited lower water sorption and lower hardness than the film containing an equal weight fraction of NVF.

Having thus described the present invention, what is now deemed appropriate for Letters Patent is set out in the following appended claims.

Claims

1. N-vinyl compositions comprising the structural formula:



where R^1 is hydrogen or methyl, and R^2 is a polycyclic alkyl, benzyl or alkylene group, each containing up to about 20 carbon atoms.

2. A composition in accordance with of Claim 1 in which R^1 is hydrogen.
3. A composition in accordance with Claim 2 in which R^2 is selected from the group consisting of: isobornyl, benzyl, allyl, N,N-dimethylaminoethyl and hexafluorobutyl.
4. A composition in accordance with Claim 2 which is isobornyl 3-(N-vinylformamido)propionate.
5. A composition in accordance with Claim 2 which is benzyl 3-(N-vinylformamido)propionate.
6. A composition in accordance with Claim 2 which is N,N-dimethylaminoethyl 3-(N-vinylformamido)propionate.
7. A composition in accordance with Claim 2 which is 2,2,3,4,4,4-hexafluorobutyl 3-(N-vinylformamido)propionate.
8. A composition in accordance with Claim 2 which is allyl 3-(N-vinylformamido)propionate.
9. A composition consisting of homopolymerized units of the structure of Claim 1.
10. A composition consisting of homopolymerized units of the structure of Claim 2.
11. A composition consisting of polymerized units of the structure of Claim 1 and a second polymerizable monomer.
12. A composition consisting of polymerized units of the structure of Claim 2 and a second polymerizable monomer.
13. The composition of Claim 11 in which the second polymerizable monomer is selected from the group consisting of: vinyl esters, vinylamides, acrylic esters, methacrylic eaters, maleic esters, substituted and unsubstituted acrylamides, substituted and unsubstituted methacrylamides, diallyl monomers; and maleic acid, maleic anhydride, ethylene, vinyl chloride, acrylonitrile, cationic acid and acrylic acid.
14. The composition of Claim 11 in which the second polymerizable monomer is a water-soluble vinylamide.
15. The composition of Claim 14 in which the vinylamide is N-vinylformamide or N-vinyl-2-pyrrolidone.
16. The composition of Claim 13 in which the second polymerizable monomer is butyl acrylate or vinyl acetate.



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EUROPEAN SEARCH REPORT

Application Number
EP 96 11 4552

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Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.6)
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A	FR-A-2 384 008 (GAF CORP) * page 7 - page 8 *	1,11	
D	& US-A-4 129 709		
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A	TETRAHEDRON LETTERS, vol. 29, no. 23, 1988, OXFORD GB, pages 2797-2780, XP002020765 L. BECKING AND H.J. SCHÄFER : "pyrrolidines by intramolecular addition of kolbe radicals..." * page 2798 *	1	
			TECHNICAL FIELDS SEARCHED (Int.Cl.6)
			C07C C08F
The present search report has been drawn up for all claims			
Place of search BERLIN		Date of completion of the search 10 December 1996	Examiner Rufet, J
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